

Atmospheric mercury concentrations near Salmon Falls Creek Reservoir - Where is the mercury coming from?

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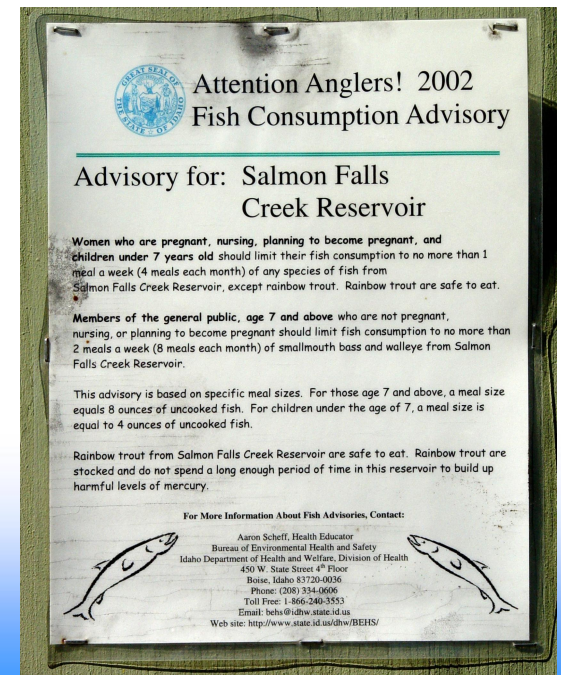


Four lakes in southern Idaho w/ Hg advisories

Salmon Falls Creek Reservoir

Fish Species	Mercury Range (mg/kg)	No. of Fish	% above 0.3
Kokanee	0.17 - 0.25	2	0
Rainbow Trout	0.0 - 0.24	11	0
Smallmouth Bass	0.44 - 0.66	2	100
Walleye	0.25 - 1.08	12	92
Yellow Perch	0.2 - 0.48	10	80

- IDEQ began study at SFCR in 2005 – air, water, snow, wet dep, sediment
- What's causing high fish concentrations?
- Our objective – is there an air input and where is it coming from?

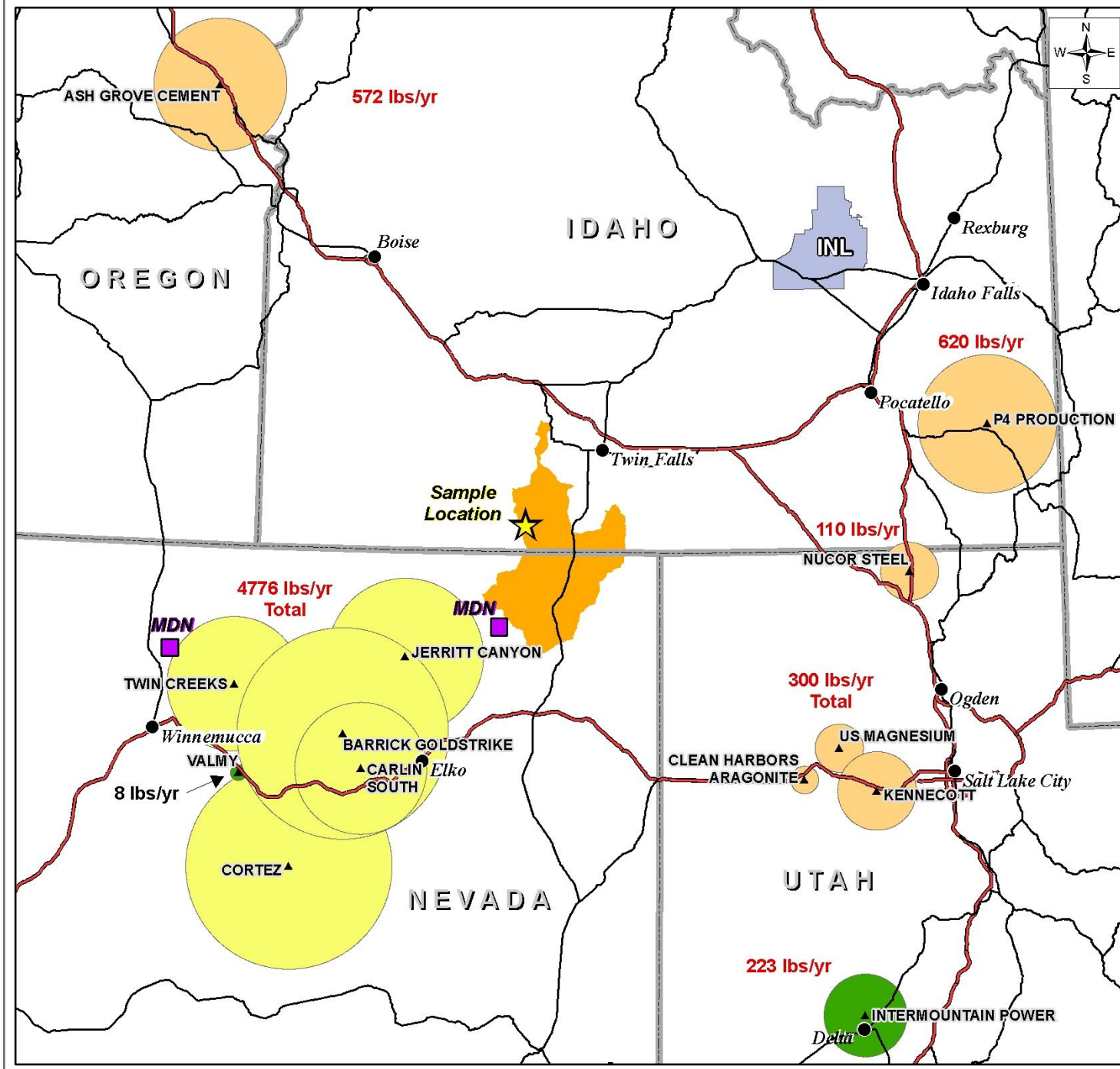


Air Input?

- Elemental gaseous Hg (EGM) - most (99%) of atmos Hg
 - relatively non-reactive, low deposition rate, but some studies suggest it may contribute.
- Reactive gaseous Hg (RGM) and particulate Hg (HgP) - oxidized Hg (mostly Hg^{+2}) - highly reactive and depositing.
 - found locally near sources with RGM/HgP emissions.
 - produced in atmos from O_3 and OH oxidation of EGM.
- RGM Wet deposition - MDN-type sampler
 - Two NV sites show sporadic high concentrations in summer (50 to >100 ng/L) , but total annual flux not unusually high ($3\text{-}4$ ug/m²/y).
- RGM Dry deposition – likely the driver for arid regions
 - no proven direct measurement methods.
 - Estimate using RGM air conc x published deposition velocities.

Why measure Hg⁰?

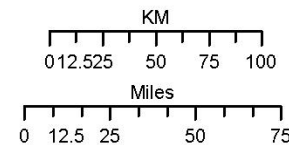
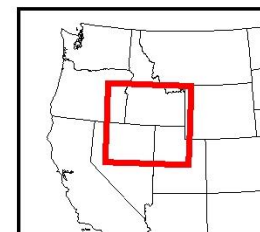
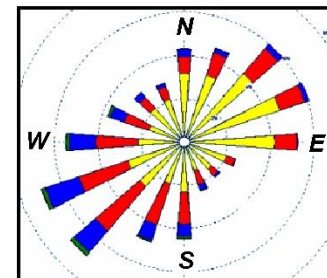
- Prevailing winds in SFCR watershed from NV; most sources in NV, both natural and anthro, emit primarily Hg⁰:
 - Evasion from naturally enriched substrates, hydrothermal areas – est. 10,000 kg/y over all of NV (Zenner and Gustin, 2002)
 - Largest anthro sources in NV are gold mines (11) - mostly Hg⁰ – 2003 est. 2,200 kg/y; 1998-2003 avg 5,100 kg/y (EPA TRI).
- A major source of Hg⁺² in remote regions is atmospheric oxidation of Hg⁰, esp. in summer when photochemical oxidation rates are high (ozone, sunlight).
- For a given photochemical oxidation rate (RGM/EGM ratio), high Hg⁰ will increase local production and deposition of Hg⁺².
- Jury still out on EGM deposition – long-term flux chamber measurements needed.



Relative Source Strength Hg lbs/yr (2003 TRI)

Legend

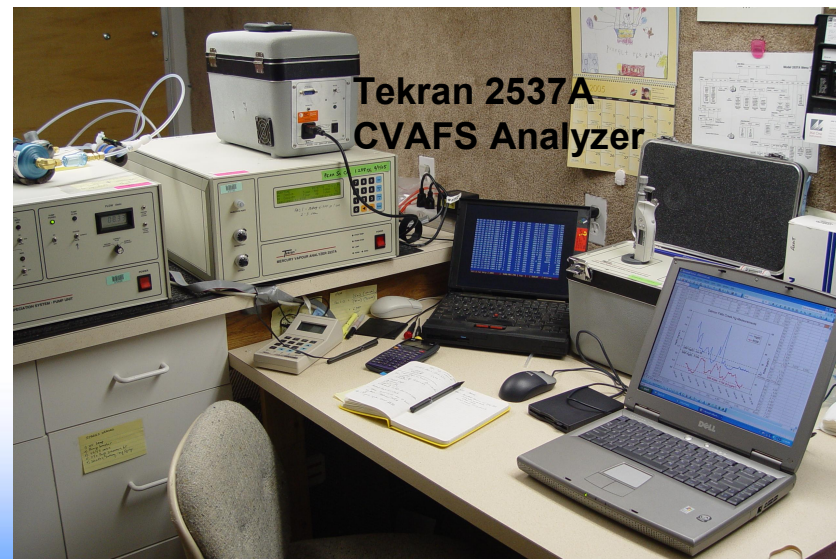
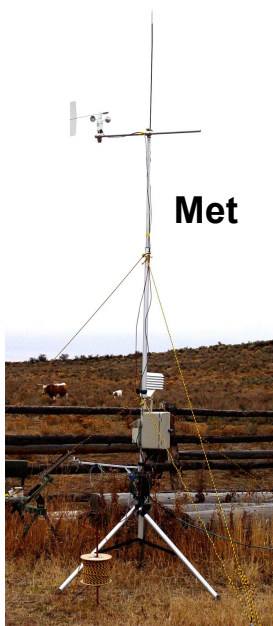
- ▲ Source Locations
- Coal-fired Power Plant
- Gold Mine
- Other Source
- Cities and Towns
- Salmon Falls Watershe
- State Boundary
- Interstate
- State/US Highway



This is a remote area!
- no local Hg sources
(nothing but cow dogs on the
lookout for coyotes)

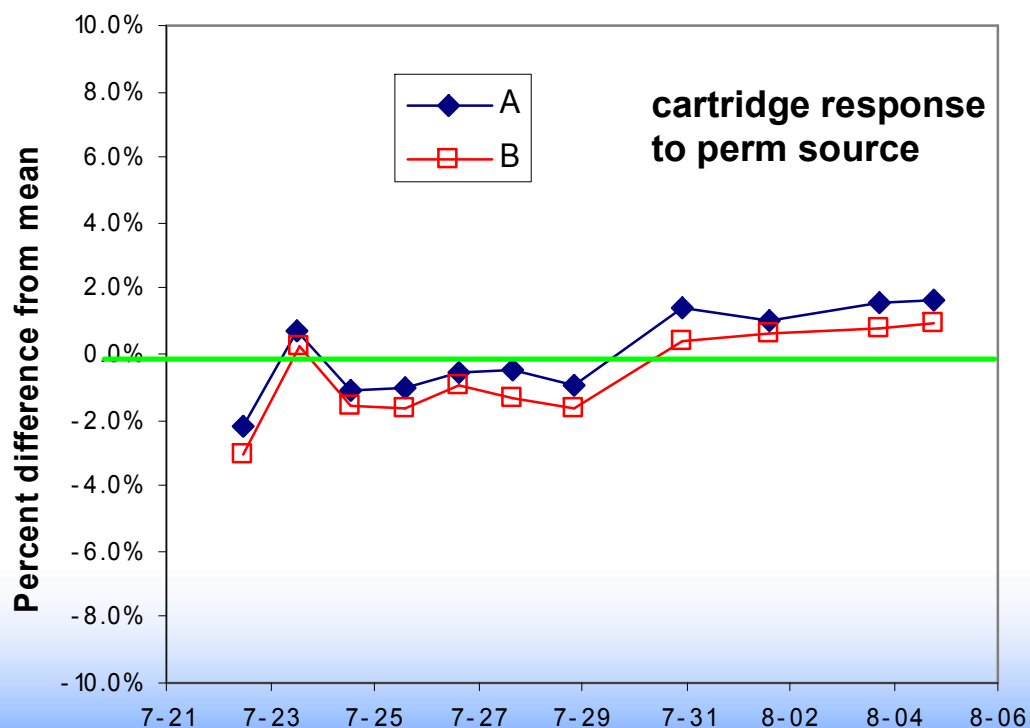


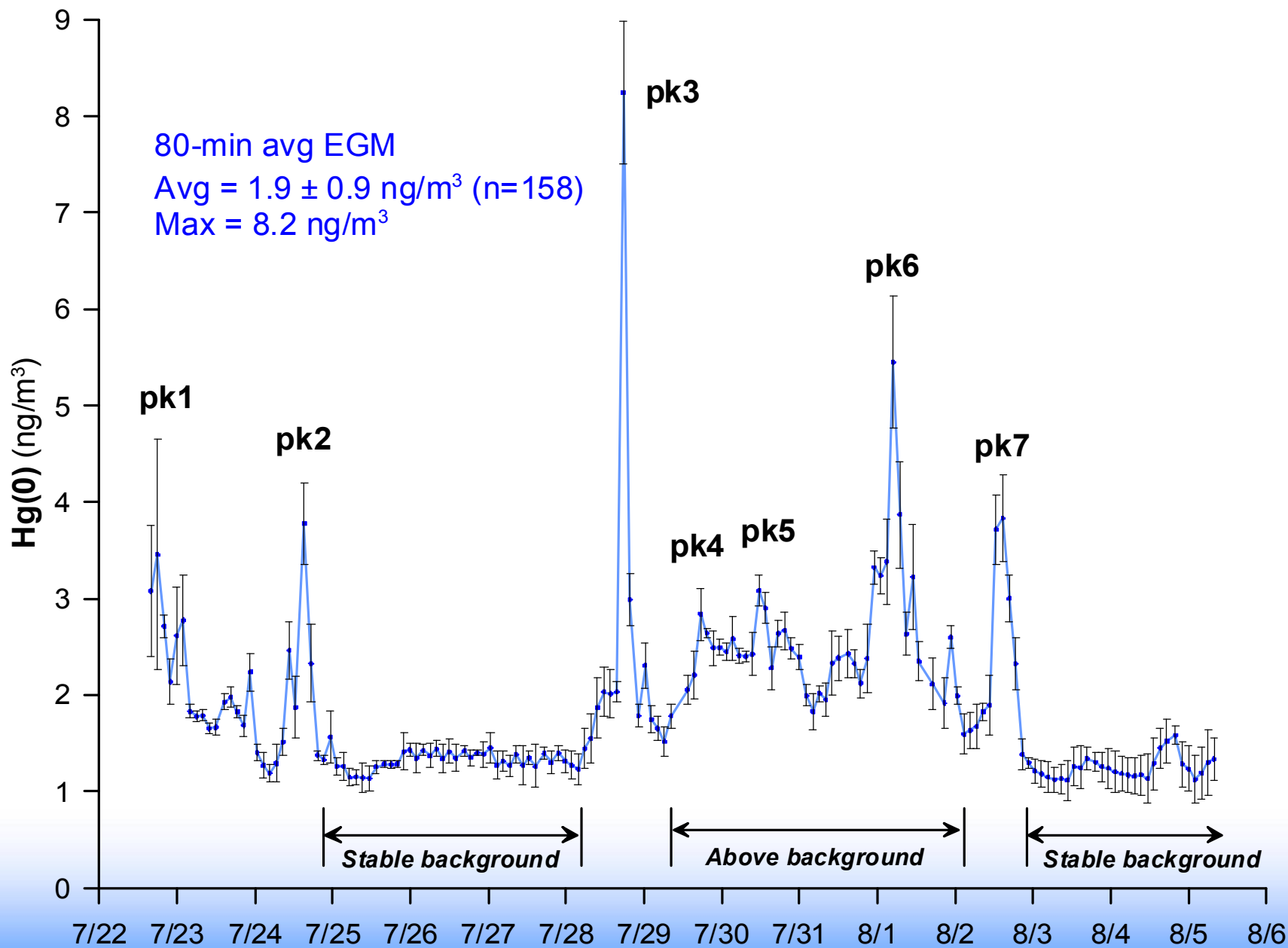
- Speciated EGM, RGM measured w/ INL Tekran 2537A, 1130, 14 day data sets-Jul/Aug and Nov 2005 (in progress)
- NO₂, SO₂, PM-10, and ozone w/ IDEQ instruments
- Met – WS,WD,AT,RH,SR



QA/QC

- 6-month Calibration of 2537A using 2505 saturated vapor calib unit (detection limit $< 0.1 \text{ ng/m}^3$).
- Standard additions ($n=12$) prior to deployment ($102\% \pm 6\%$)
- 2537A mass flow controller calibration ($+2\%$)
- Total system blank check w/ zero air (always 0)
- Total system standard additions w/ vapor injections ($95\text{-}110\%$)
- Daily perm source calibrations ($\pm 1.3\%$)
- Cartridge bias check

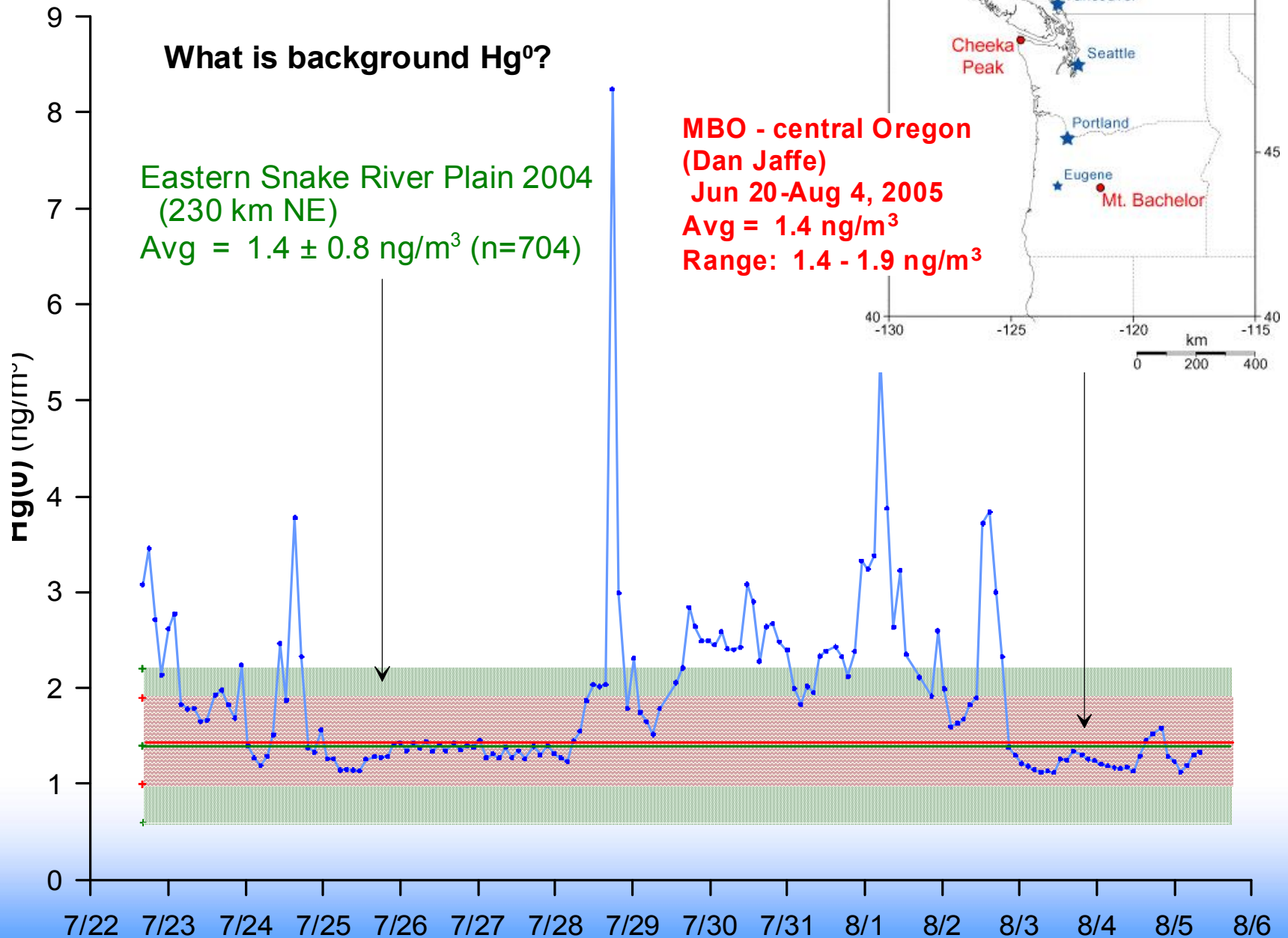


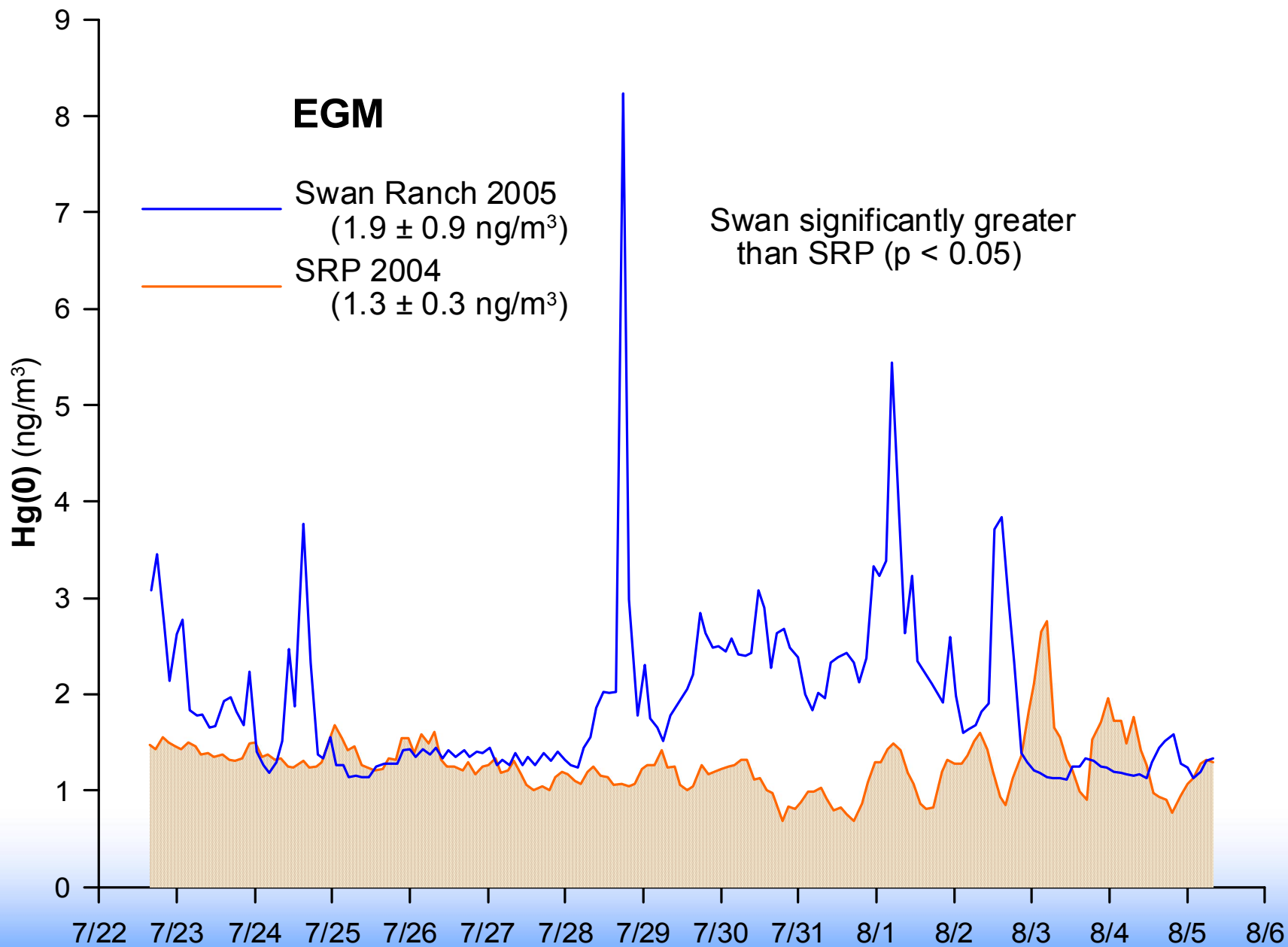


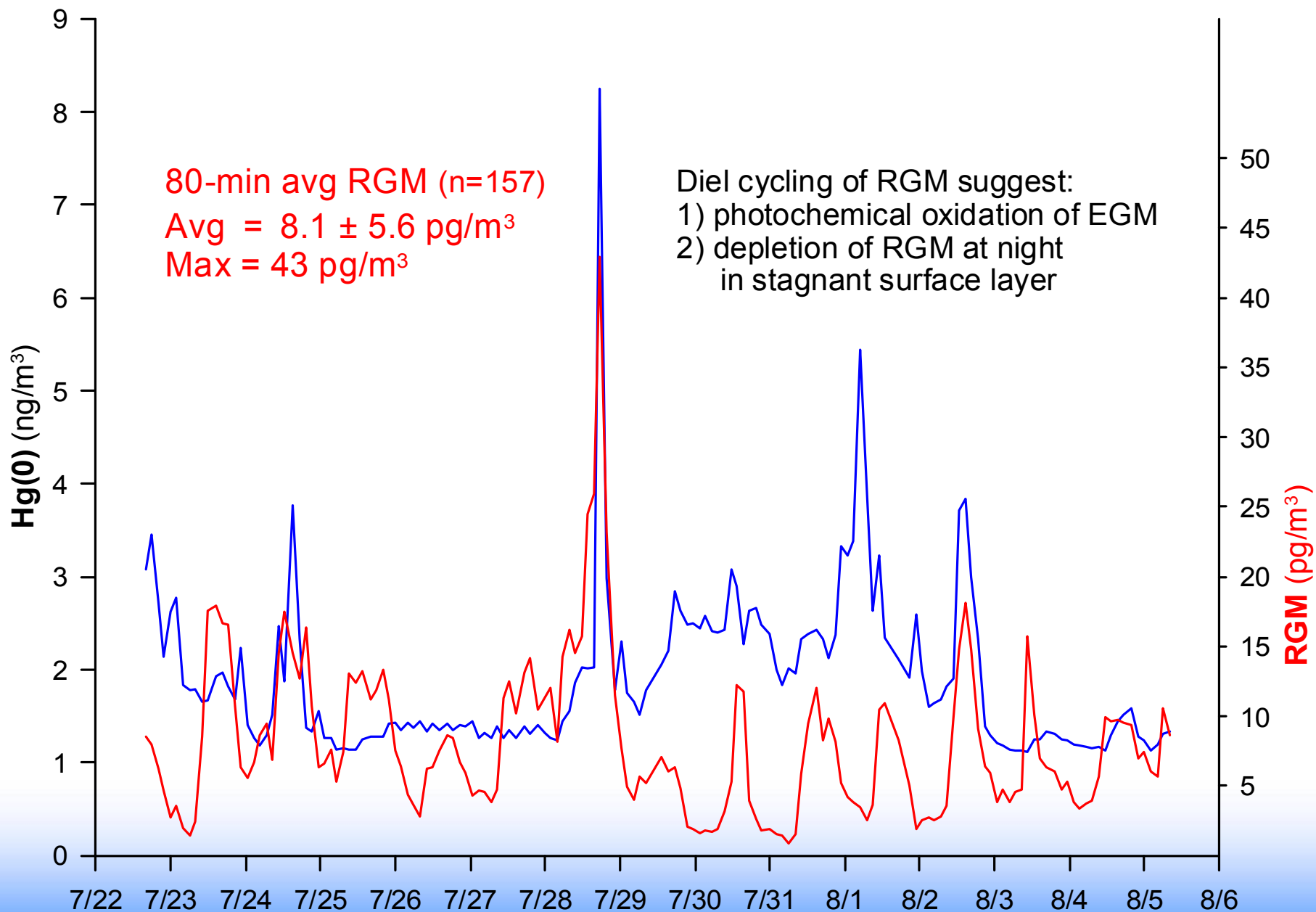
What is background Hg⁰?

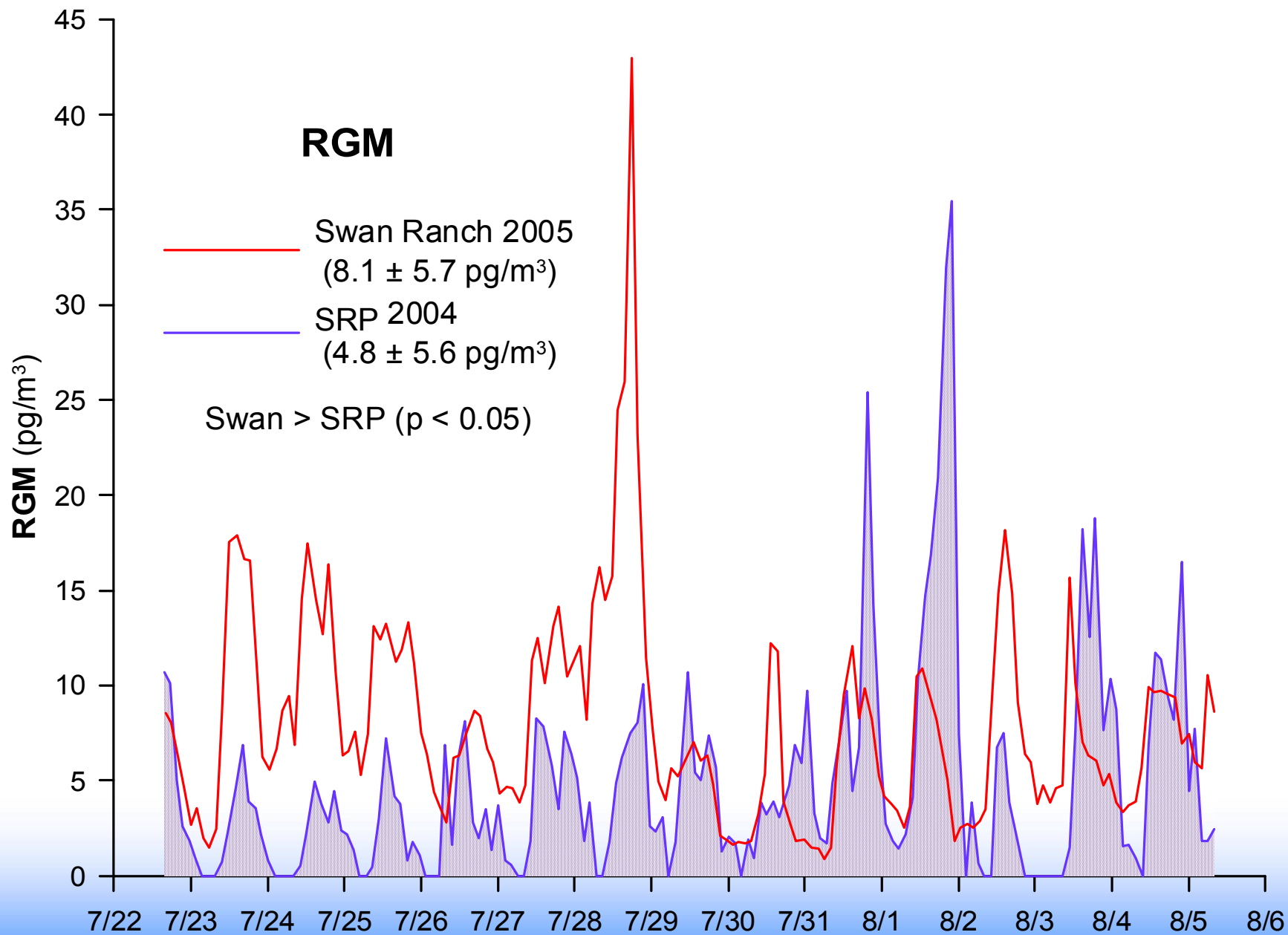
Eastern Snake River Plain 2004
(230 km NE)
Avg = $1.4 \pm 0.8 \text{ ng/m}^3$ (n=704)

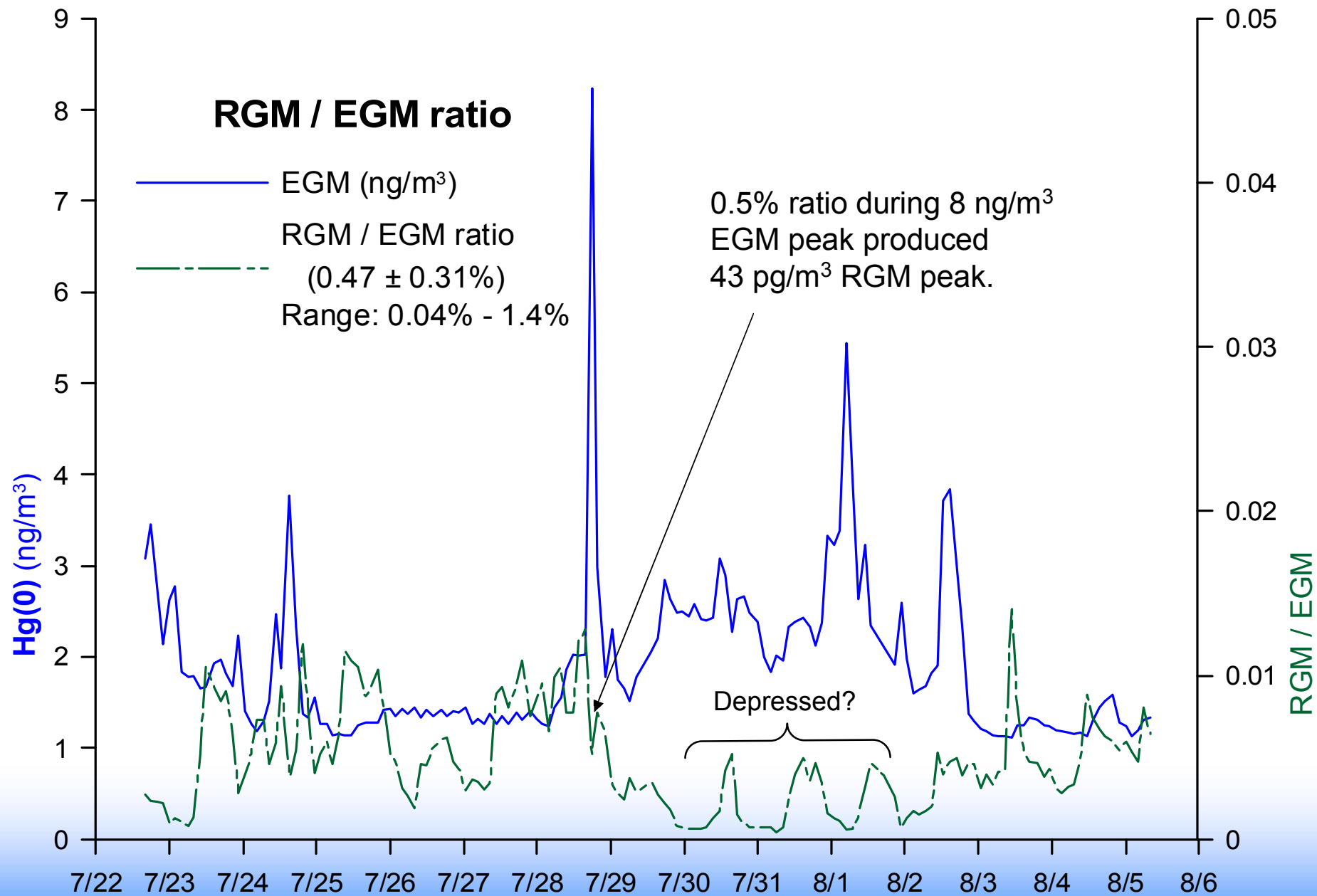
**MBO - central Oregon
(Dan Jaffe)
Jun 20-Aug 4, 2005
Avg = 1.4 ng/m^3
Range: $1.4 - 1.9 \text{ ng/m}^3$**



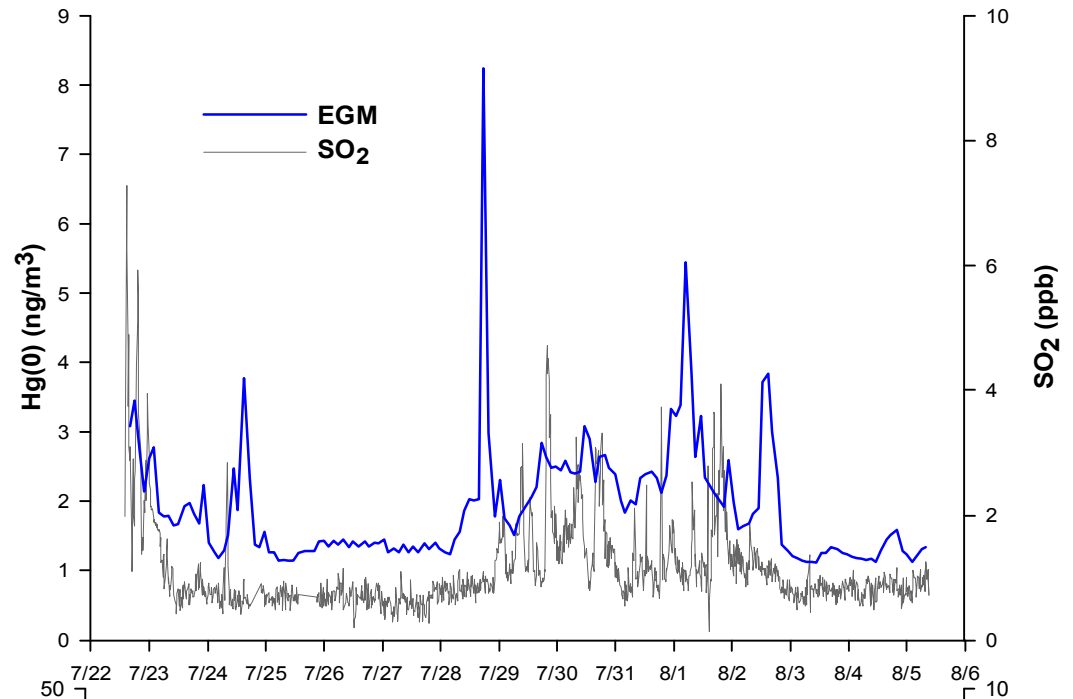




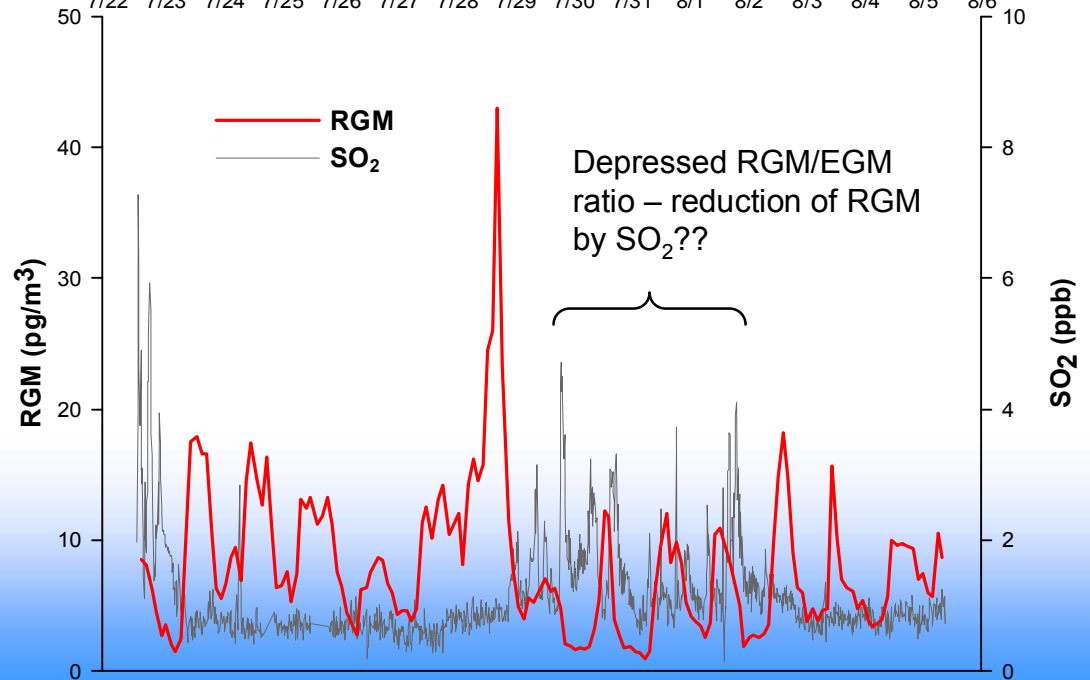




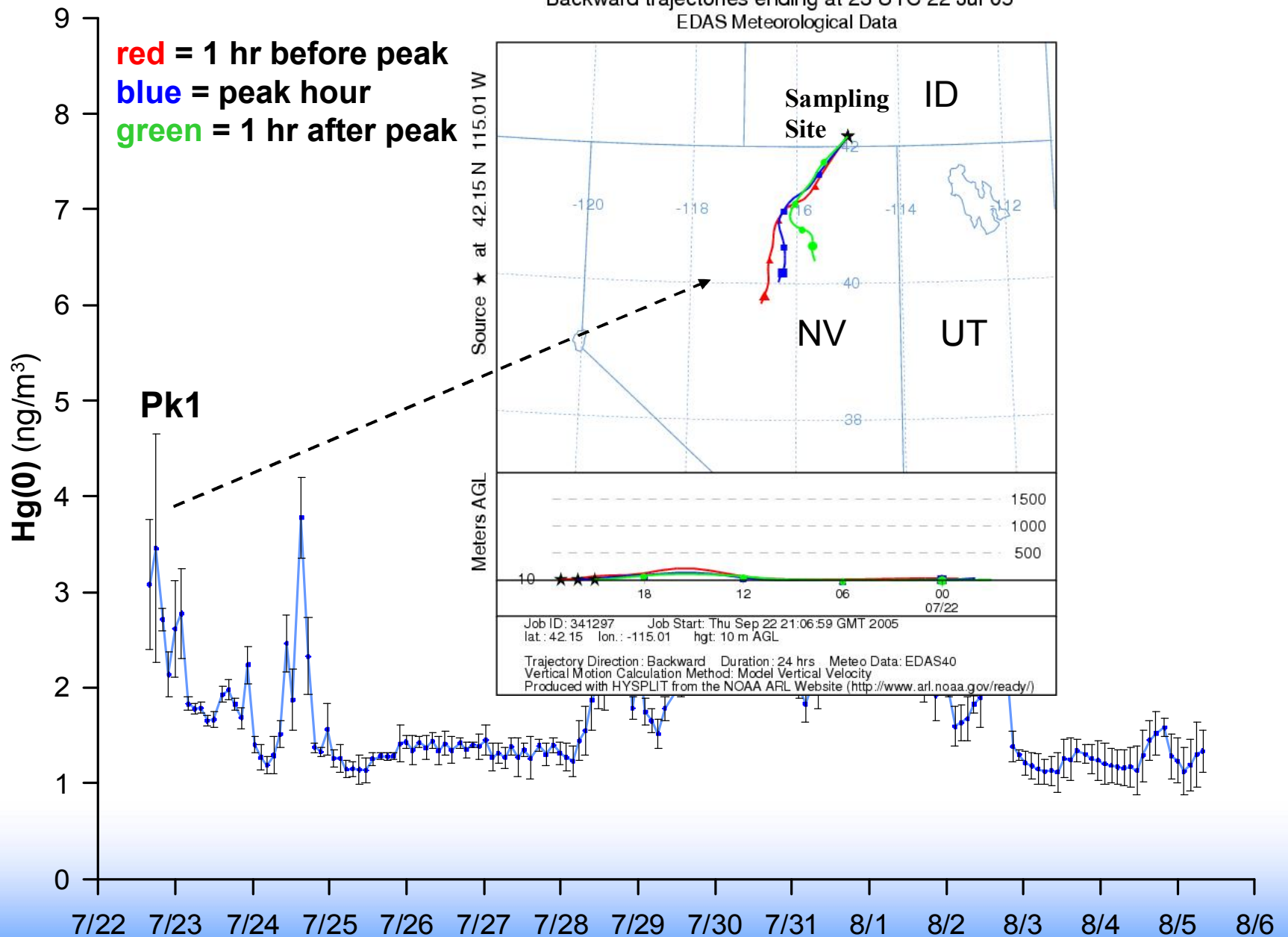
EGM and SO₂



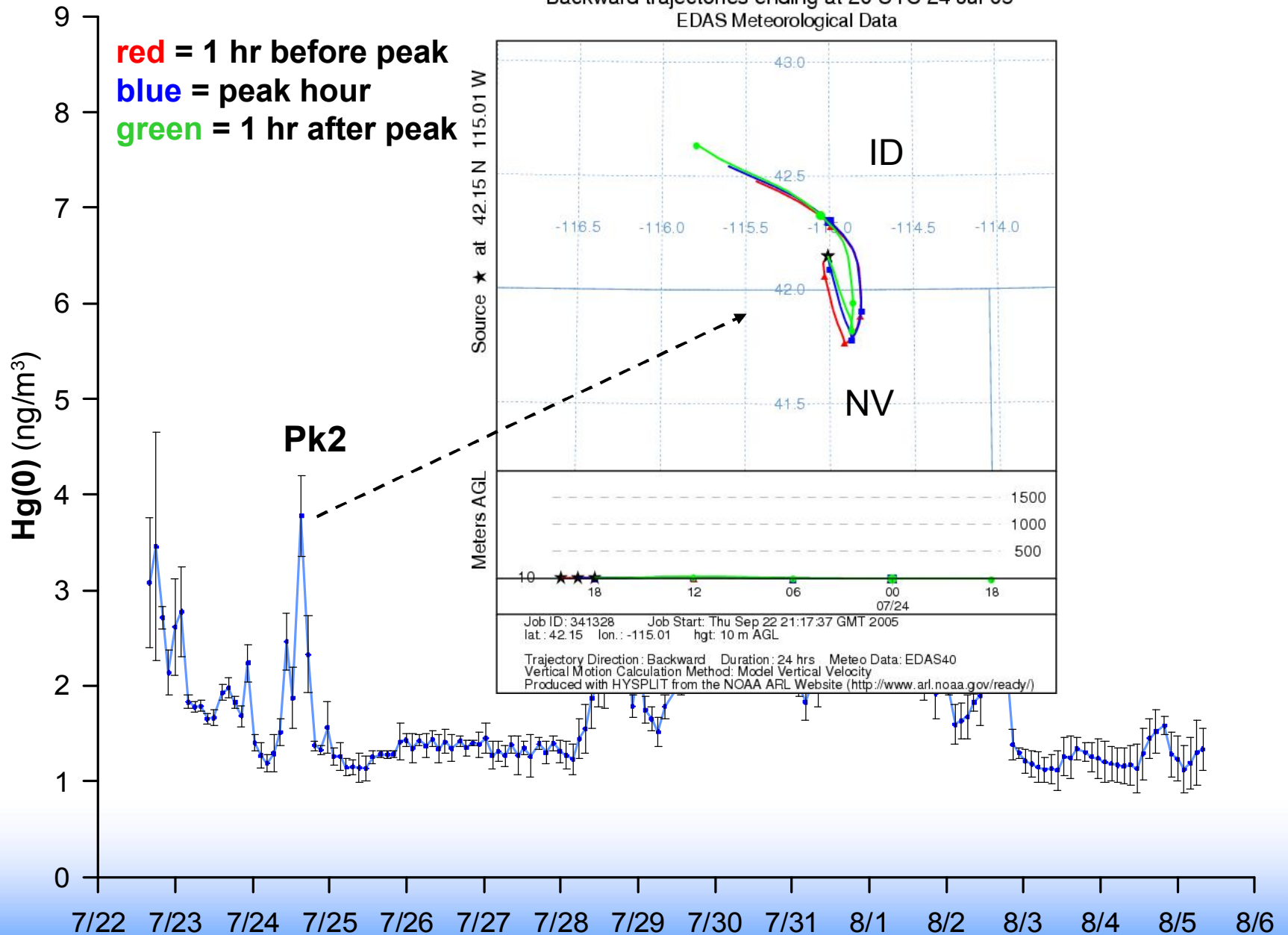
RGM and SO₂



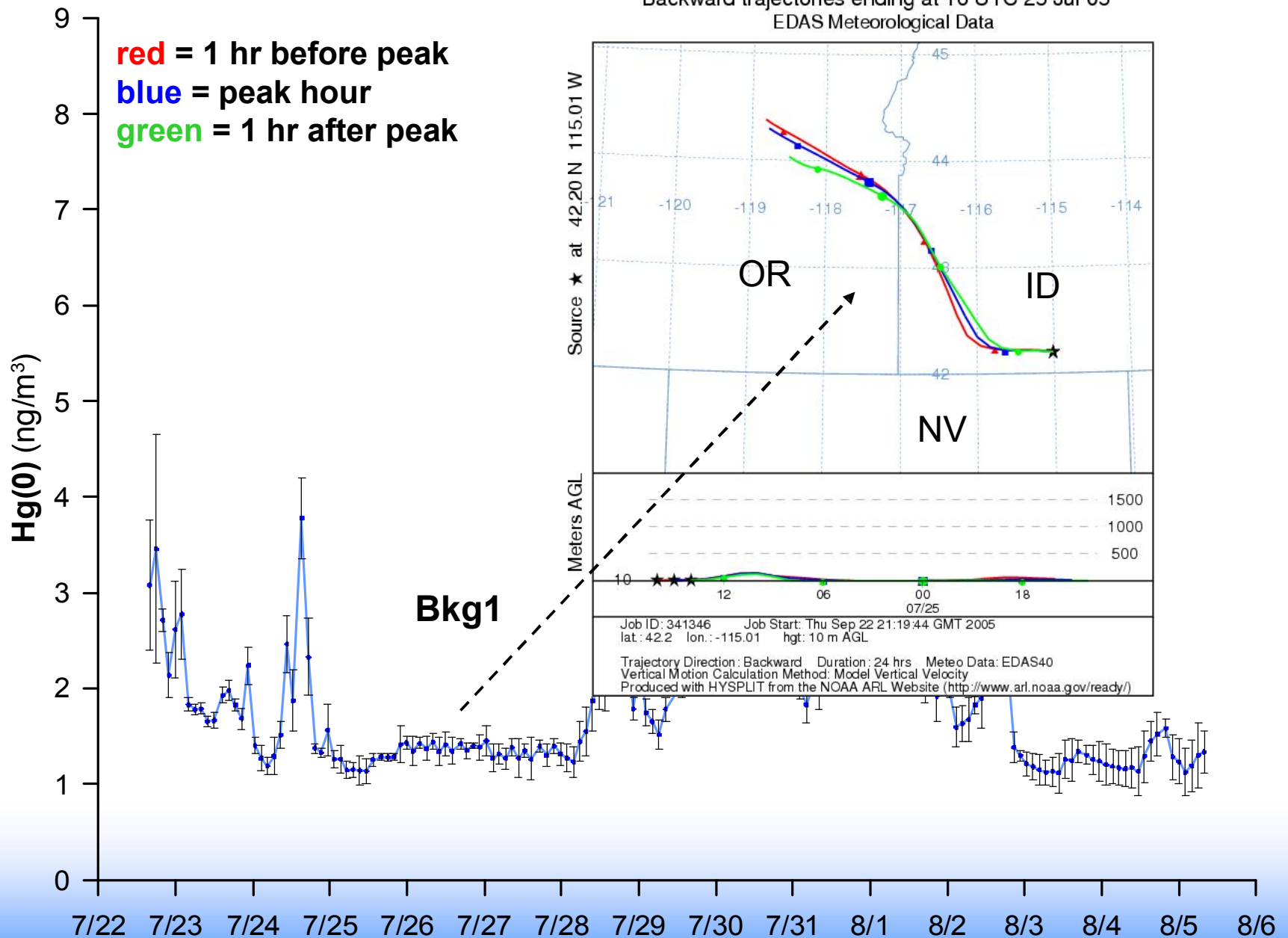
NOAA HYSPLIT MODEL
Backward trajectories ending at 23 UTC 22 Jul 05
EDAS Meteorological Data



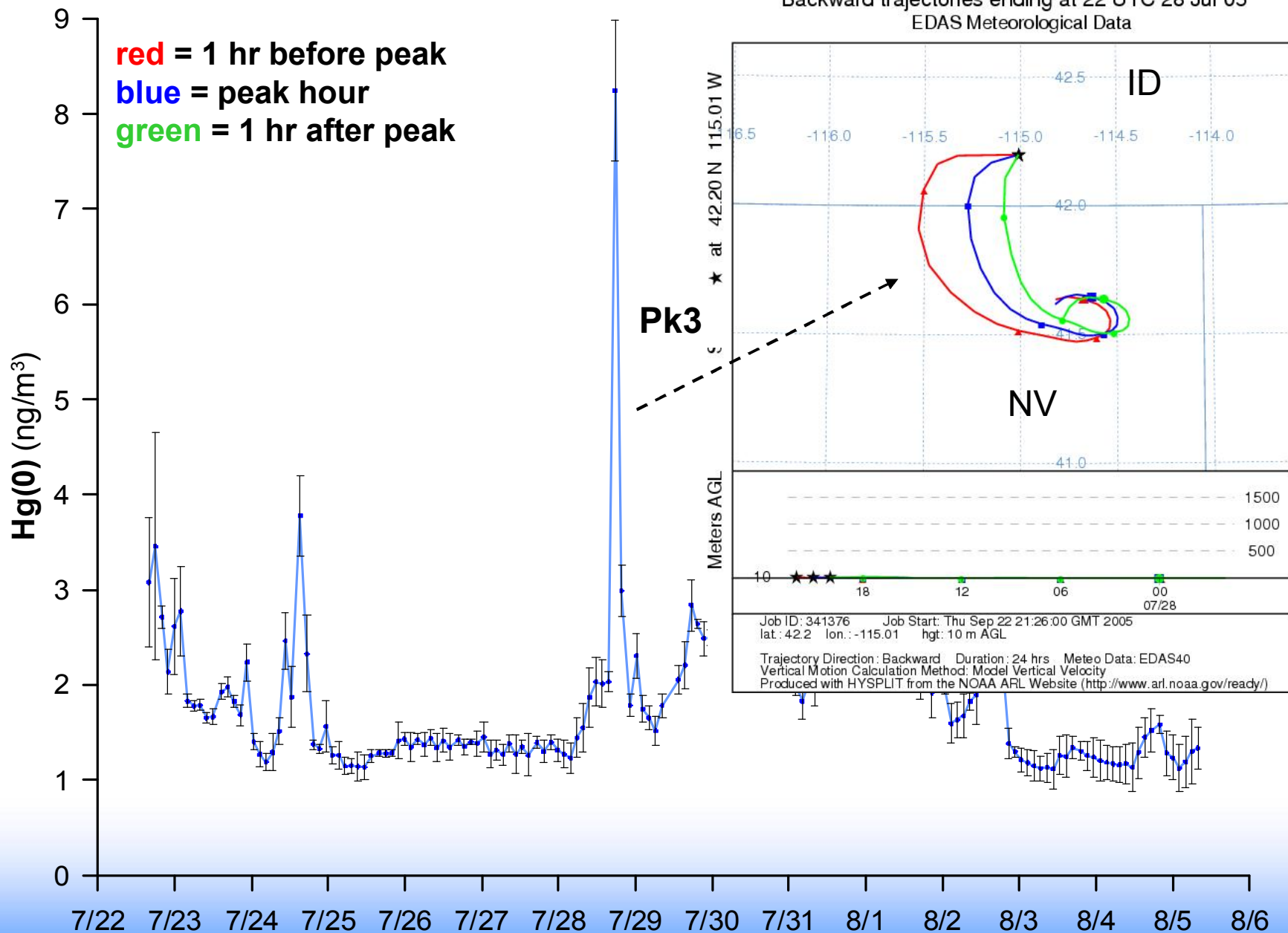
NOAA HYSPLIT MODEL
Backward trajectories ending at 20 UTC 24 Jul 05
EDAS Meteorological Data

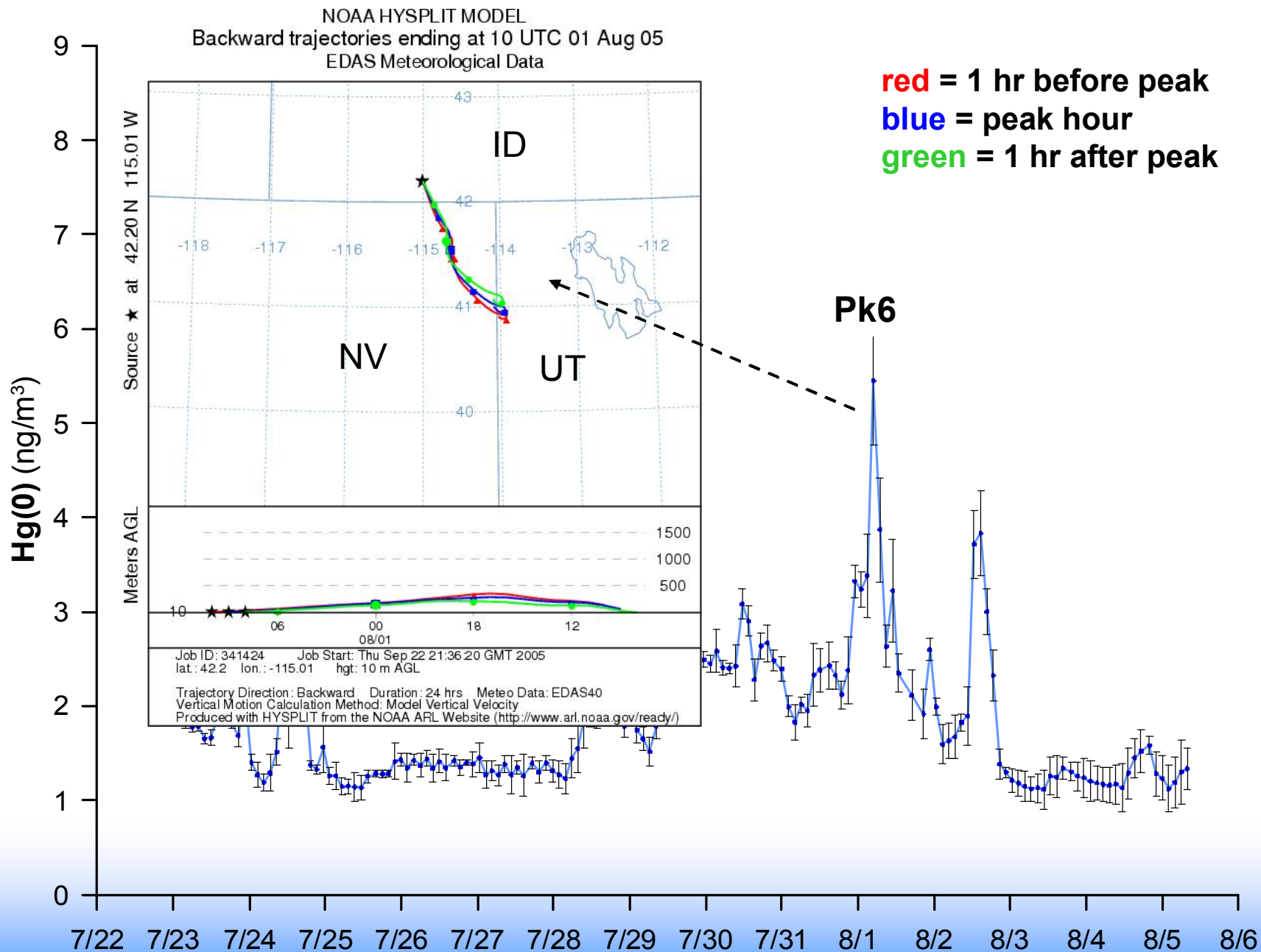


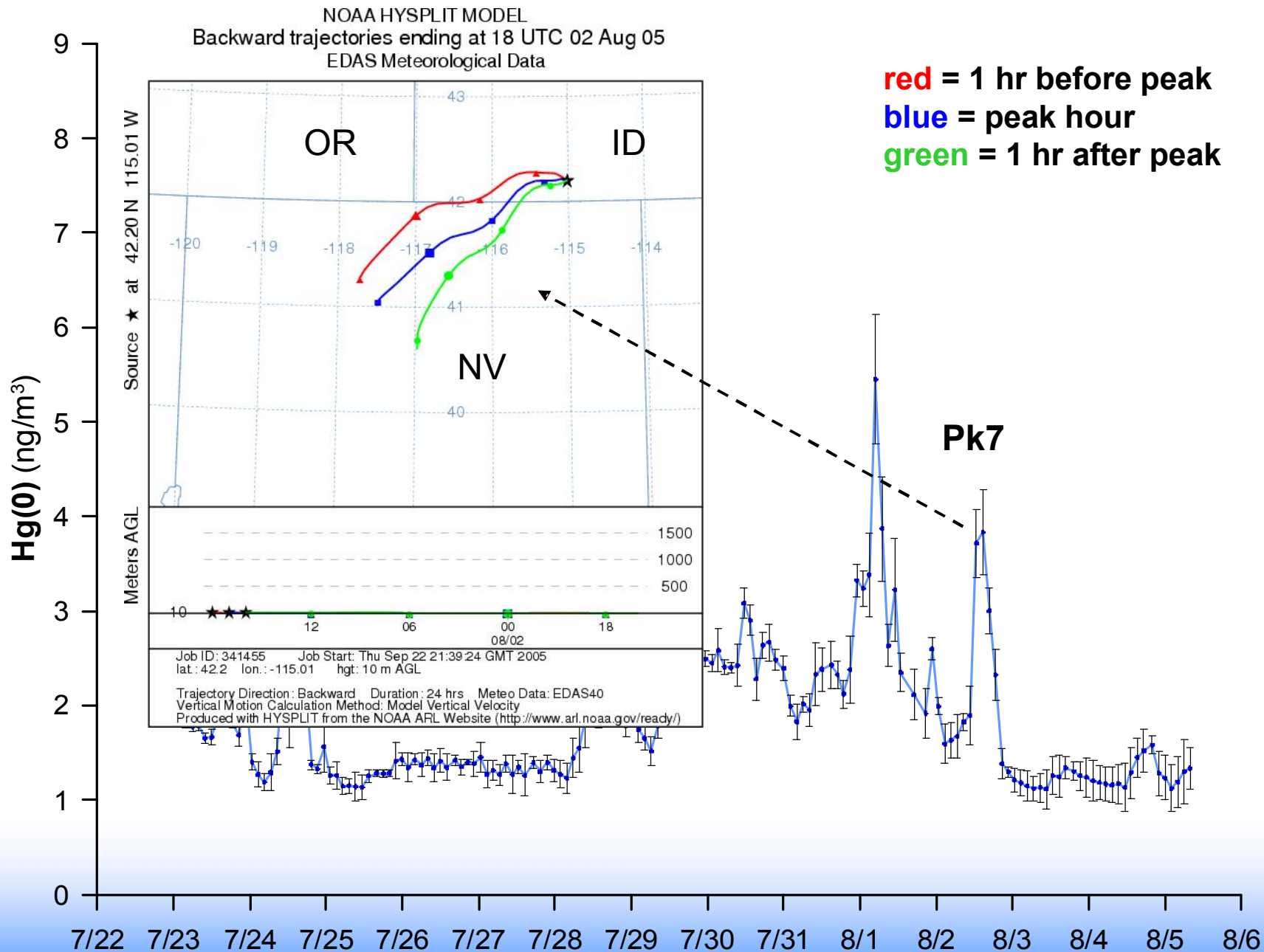
NOAA HYSPLIT MODEL
Backward trajectories ending at 16 UTC 25 Jul 05
EDAS Meteorological Data

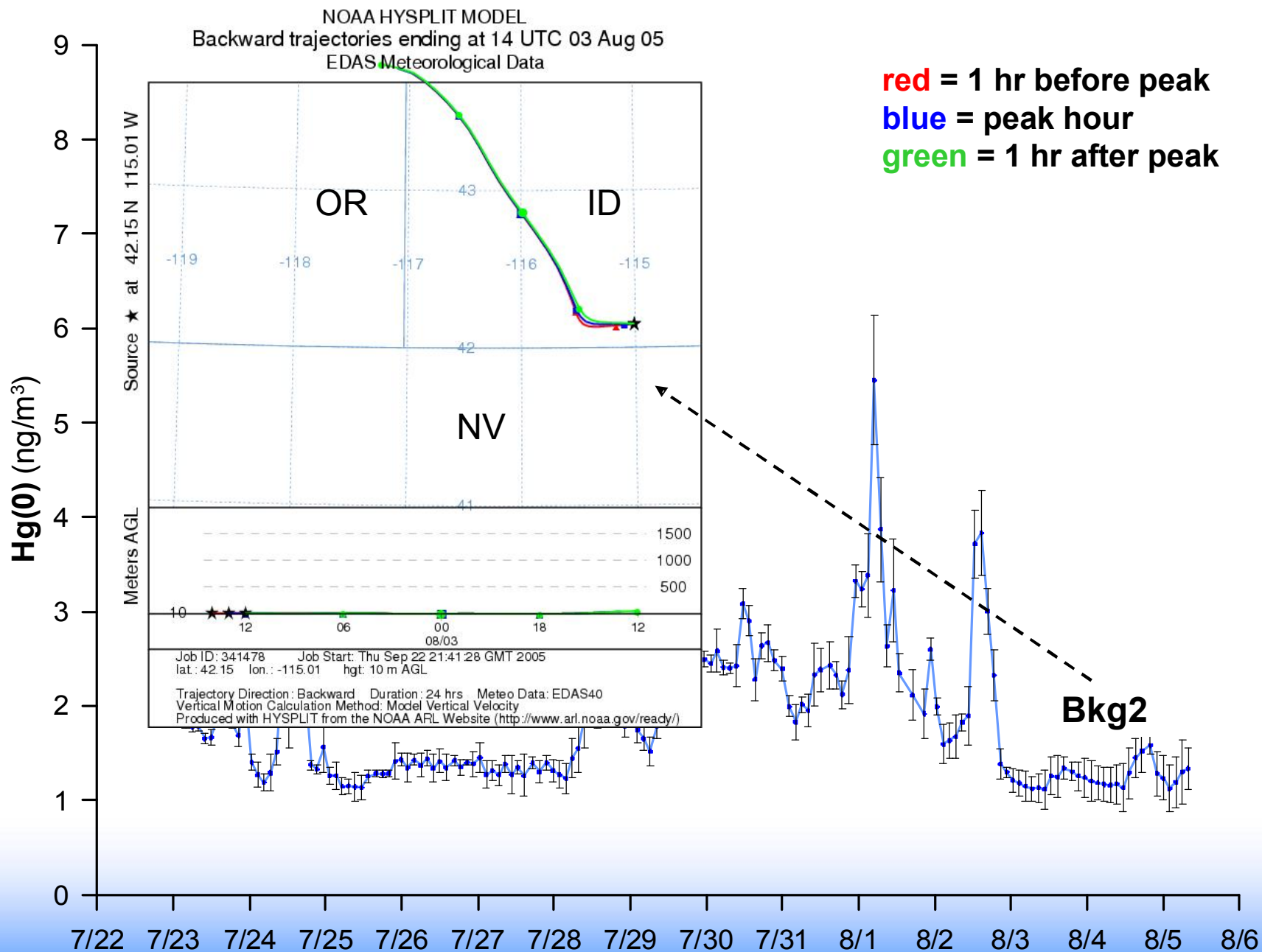


NOAA HYSPLIT MODEL
Backward trajectories ending at 22 UTC 28 Jul 05
EDAS Meteorological Data

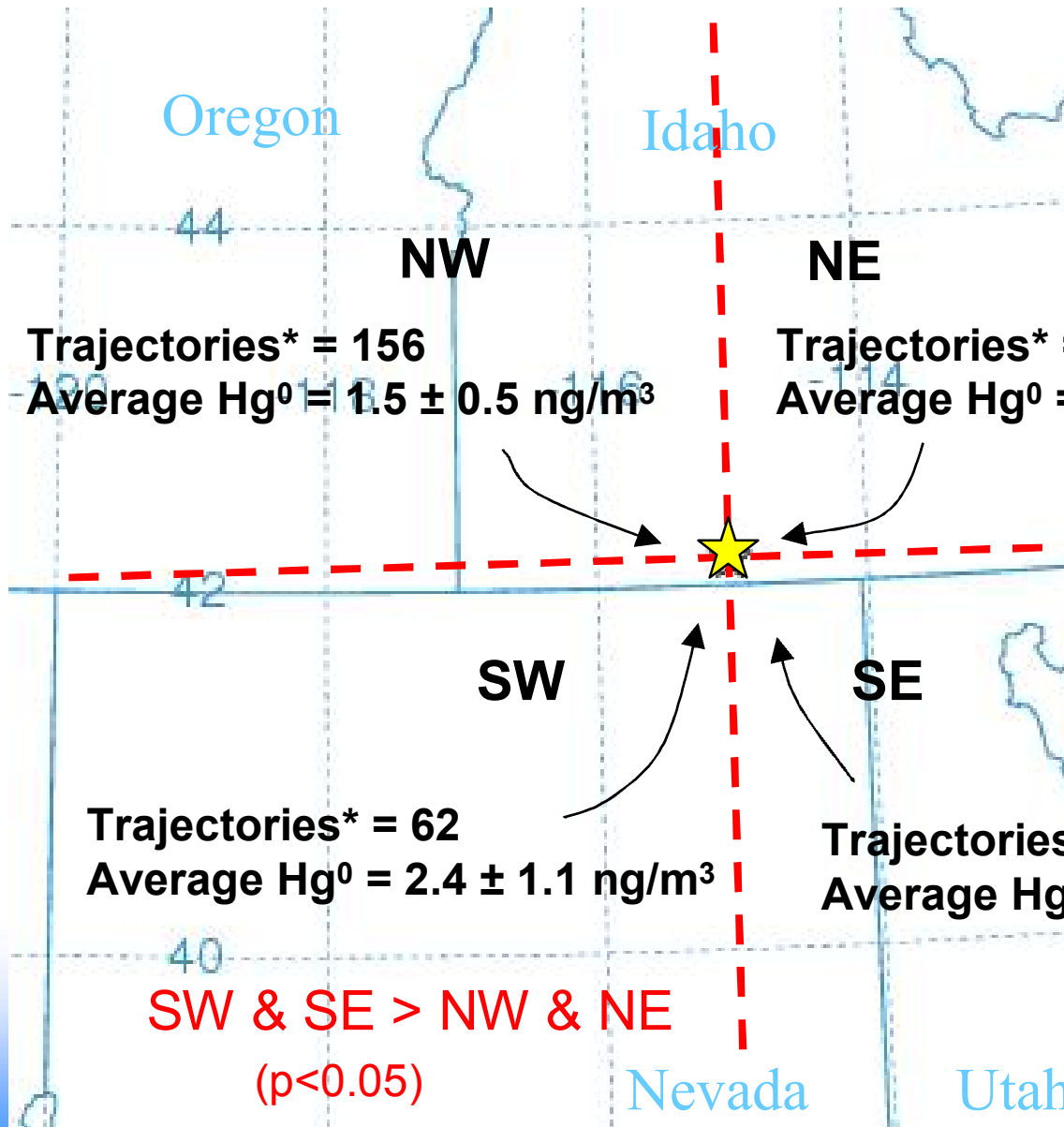








Trajectory analysis by quadrant



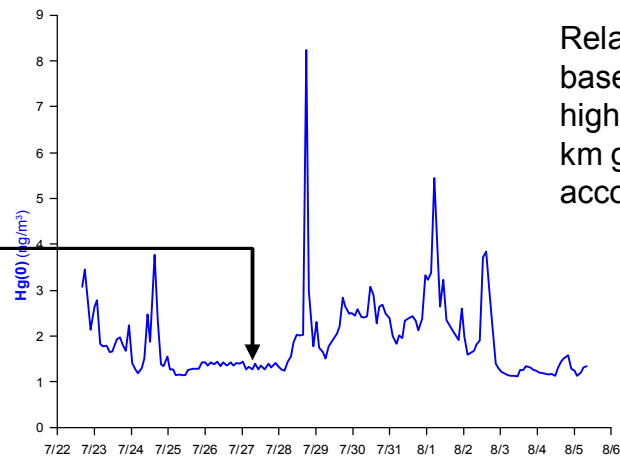
★ --- Monitoring location

*Trajectories were assigned to a quadrant based on where they spent the majority of time in the previous 24 hours.

NV Gold mine input?

NV Gold mine dispersion during observed background

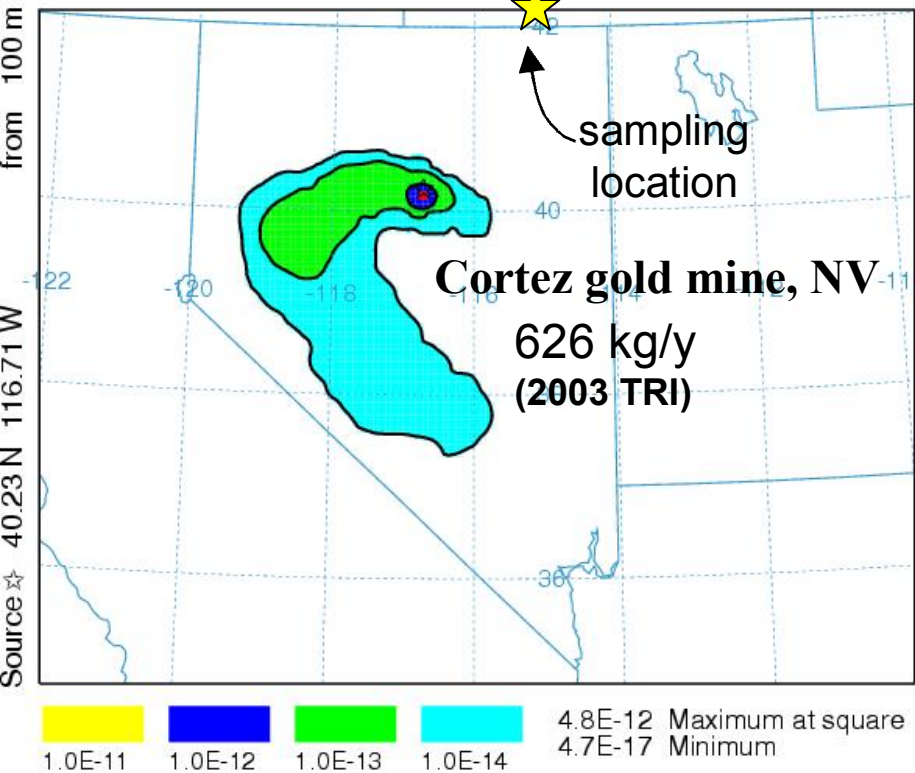
Forward trajectories:



Relative dispersion estimates based on a hypothetical 100-m high unit release. EDAS 40-km grid does not take into account local terrain.

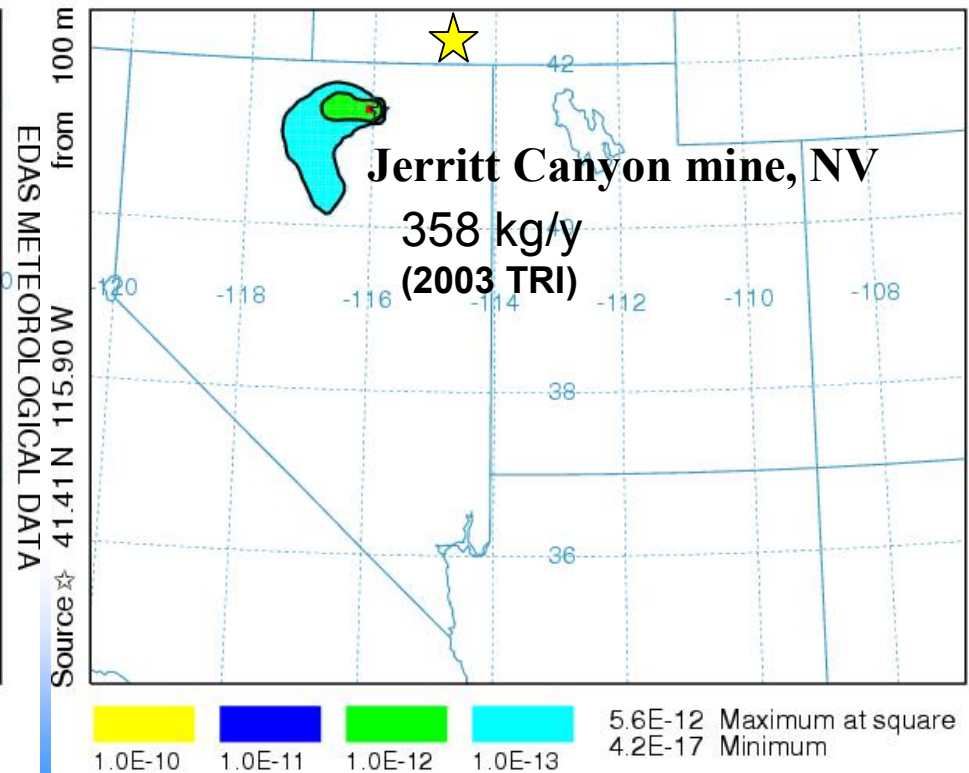
NOAA HYSPLIT MODEL

Concentration (mass/m³) averaged between 0 m and 2000 m
Integrated from 2300 26 Jul to 0000 27 Jul 05 (UTC)
Release started at 0000 26 Jul 05 (UTC)

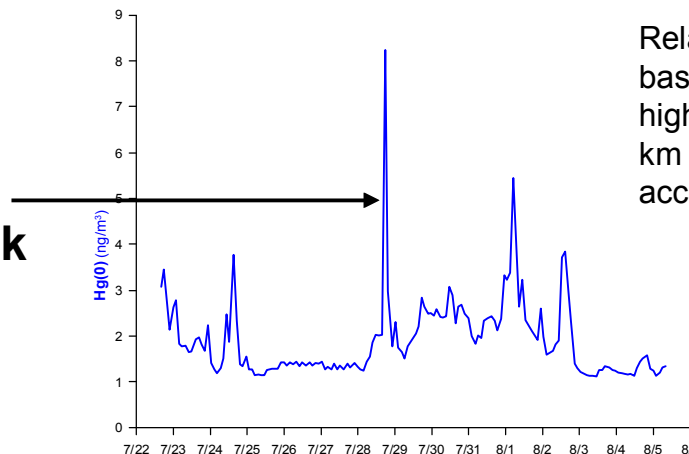


NOAA HYSPLIT MODEL

Concentration (mass/m³) averaged between 0 m and 2000 m
Integrated from 2300 26 Jul to 0000 27 Jul 05 (UTC)
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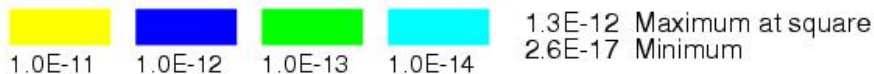
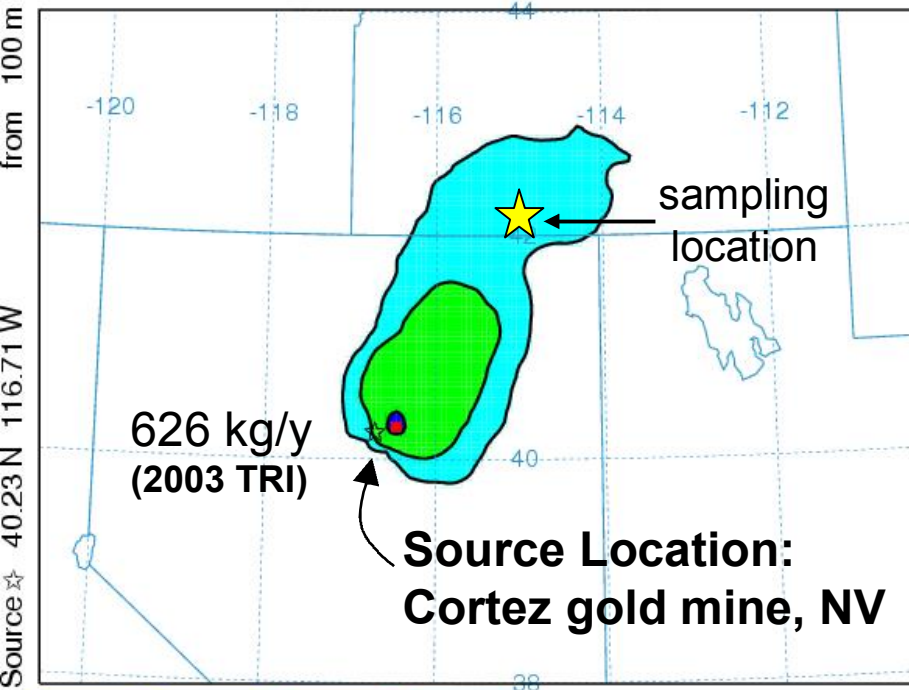
NV Gold mine dispersion during observed 8 ng/m³ peak



Relative dispersion estimates based on a hypothetical 100-m high unit release. EDAS 40-km grid does not take into account local terrain.

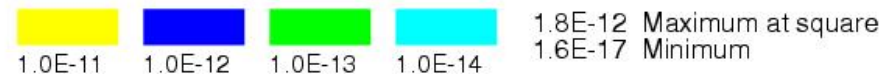
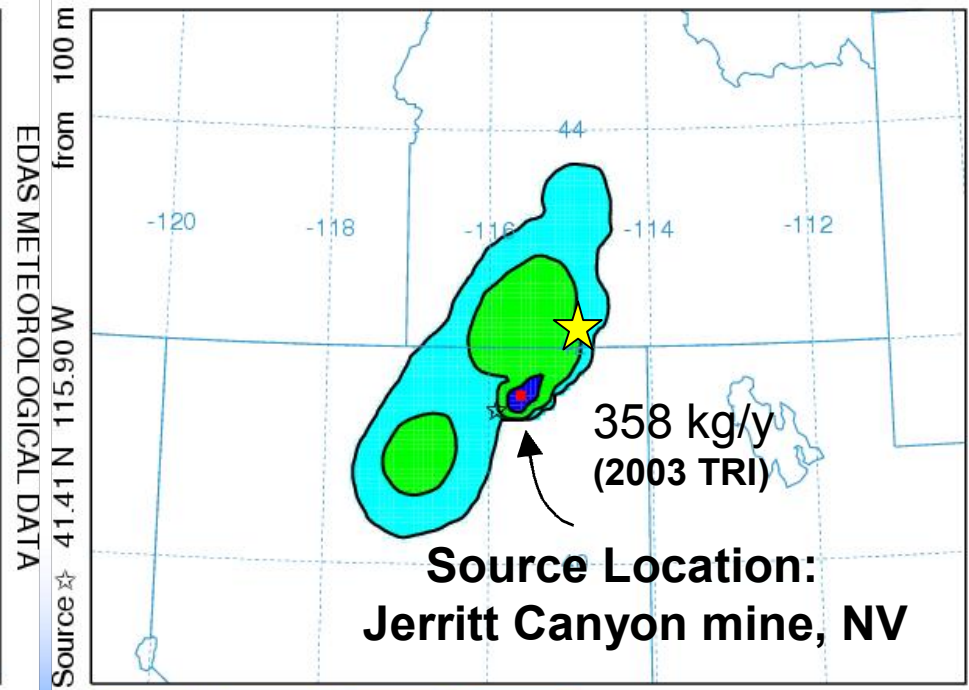
NOAA HYSPLIT MODEL

Concentration (mass/m³) averaged between 0 m and 2000 m
Integrated from 2200 28 Jul to 2300 28 Jul 05 (UTC)
Release started at 2300 27 Jul 05 (UTC)



NOAA HYSPLIT MODEL

Concentration (mass/m³) averaged between 0 m and 2000 m
Integrated from 2200 28 Jul to 2300 28 Jul 05 (UTC)
Release started at 2300 27 Jul 05 (UTC)



Natural source input?

Largest in vicinity is the Ivanhoe Mining District ~ 50 km SW of Jerritt Canyon Mine and ~ 170 km SW of the sampling location.

78 kg/y Hg emissions estimate over 582 km² undisturbed naturally enriched substrate (Engle et al. Atmos Environ 35, 2001)

Hg air concentration averaged over this area:

“box model”

$$C = \frac{\Delta x Q_A}{Z_i u} \quad (\text{Hanna et al., Handbook on Atmospheric Diffusion})$$

where

Δx = area source width ~23,000 m

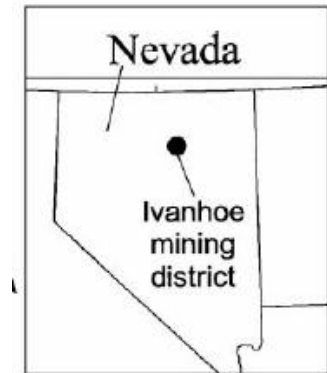
Q_A = area-averaged emission rate - 78 kg/y/582 km² = 4.3E-3 ng/m²/s ~ 15 ng/m²/hr

Z_i = mixing depth (assume 100 m)

u = annual average wind speed (assume 3 m/s)

$C = \underline{0.3 \text{ ng/m}^3}$ (fully mixed over source – rough estimate)

Dispersion to sampling site (170 km) would reduce this concentration at least 2 orders of magnitude, i.e., non-detectable.



Conclusions

- Average EGM and RGM air concentrations - significantly elevated in the SFCR area compared to background (~ 30 – 70%).
- Elevated RGM concentrations sometimes occurred with EGM peaks, likely due to atmos oxidation. Suggests EGM loading from nearby sources could indirectly increase RGM deposition.
- Elevated EGM and RGM occurred when air parcels came out of SW (Nevada) and WE (Nevada/Utah). Background concentrations occurred when the air was from the NE and NW (Oregon/Idaho).
- Most of the observed peaks were likely from NV gold mine sources--based on their back-trajectories and because these sources are the only ones large enough in the region to produce the peak concentrations observed at our sampling site.
- The relative importance of these short-term peaks for long-term watershed Hg loading (critical factor affecting fish concentrations) is not known – need to better quantify their annual frequency and magnitude compared to that from other sources.